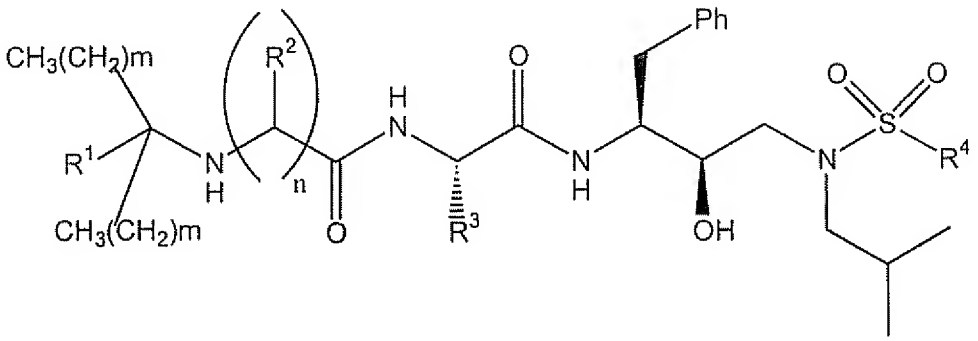
CLAIM AMENDMENTS

Please cancel claim 12.

1. (original) A compound of Formula (I)



(I)

wherein:

R^1 is phenyl, pyridinyl, pyrimidinyl, pyrazinyl, or pyridazinyl, and is unsubstituted or substituted with 1-2 substituents selected from the group consisting of halogen, C_{1-6}alkyl, trifluoromethyl, C_{1-6}alkoxy, trifluoromethoxy, and cyano;

R^2 is independently hydrogen or C_{1-6}alkyl;

R^3 is selected from the group consisting of isopropyl, tert-butyl, sec-butyl, and $C(CH_3)_2SCH_3$;

R^4 is phenyl, indazolyl, benzothiazolyl, quinolinyl, quinoxalinyl, 2,3-dihydrobenzofuranyl, or 1,3-benzodioxolyl, and is unsubstituted or substituted with 1-2 substituents selected from the group consisting of

amino, acetamido, halo, C_{1-6}alkyl, trifluoromethyl, C_{1-6}alkoxy, trifluoromethoxy, and cyano;

m is independently selected from 0, 1, 2, or 3; and

n is 1 or 2;

or a pharmaceutically acceptable salt thereof.

2. (original) A compound of claim 1 wherein R^1 is phenyl, 3-fluorophenyl, 4-fluorophenyl, 3,5-difluorophenyl, 2,4-difluorophenyl, 2,6-difluorophenyl, 3-trifluoromethylphenyl, 3-pyridinyl, or 3-methoxyphenyl.

3. (original) A compound of claim 2 where R^1 is phenyl, 3-fluorophenyl, 4-fluorophenyl, 3,5-difluorophenyl, or 3-methoxyphenyl.

4. (original) A compound of claim 1 where R^2 is hydrogen or methyl.

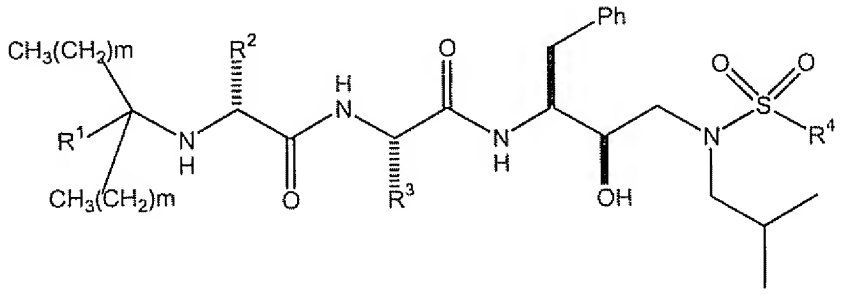
5. (original) A compound according to Claim 1 wherein R^4 is selected from the group consisting of 4-aminophenyl, 3-aminophenyl, 3-amino-4-methylphenyl, 4-methoxyphenyl, 6-benzothiazolyl, 2-amino-6-benzothiazolyl, 2-acetamido-6-benzothiazolyl, 2-methyl-6-benzothiazolyl, 7-benzothiazolyl, 2-amino-7-benzothiazolyl, 2-acetamido-7-benzothiazolyl, 2-methyl-7-benzothiazolyl, 2,3-dihydrobenzofuran-5-yl, 2,3-benzodioxl-5-yl, 6-indazolyl, 6-quinolinyl, and 6-quinoxalinyl.

6. (original) The compound according to Claim 5 wherein R^4 is selected from 4-aminophenyl, 6-indazolyl, 6-benzothiazolyl, 2-amino-6-benzothiazolyl, 6-quinolinyl, and 4-methyl-3-aminophenyl.

7. (original) A compound according to Claim 1 wherein m is 0.

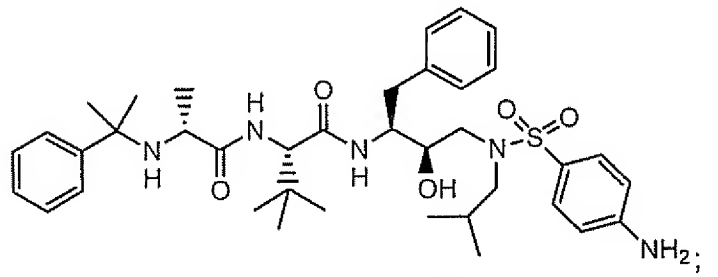
8. (original) A compound according to Claim 1 wherein n is 1.

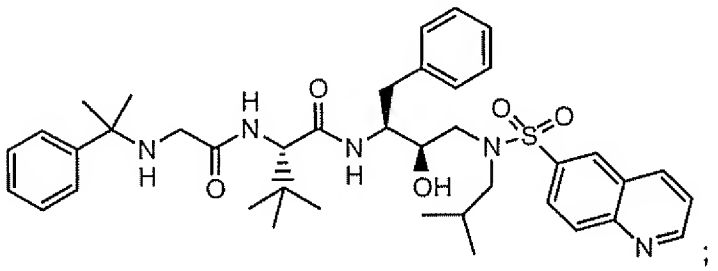
9. (original) A compound of claim 1 according to Formula Ia.

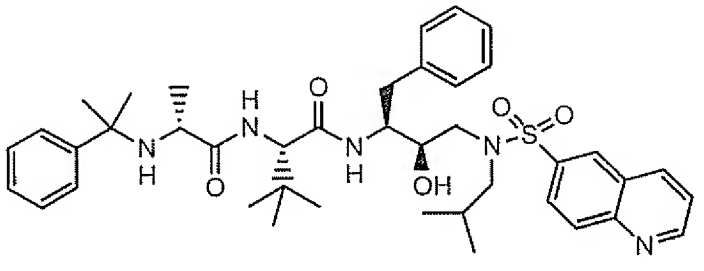


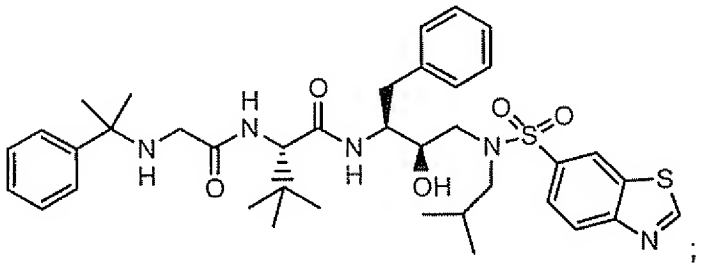
Ia

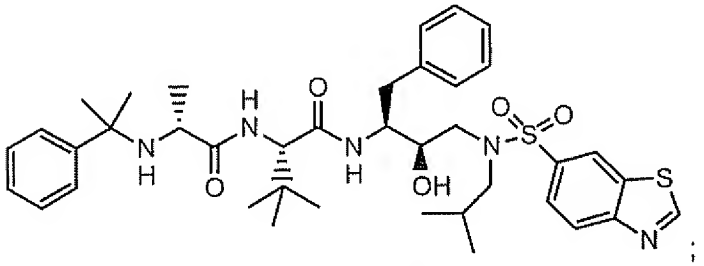
10.(original) A compound of Claim 1 selected from the group consisting of

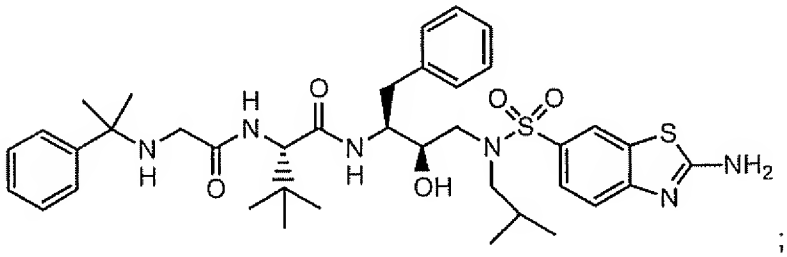


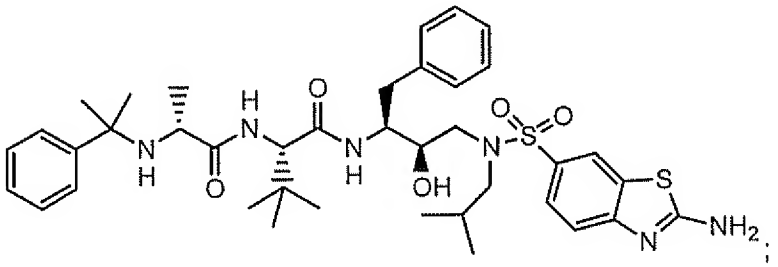


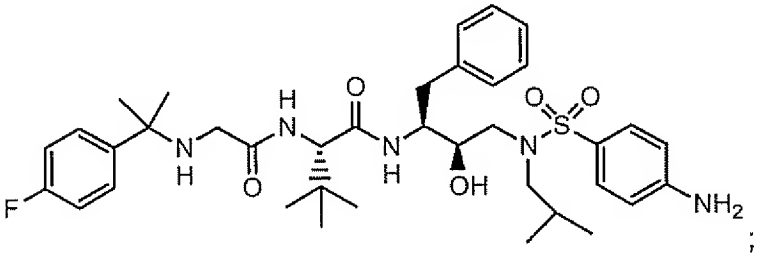

;

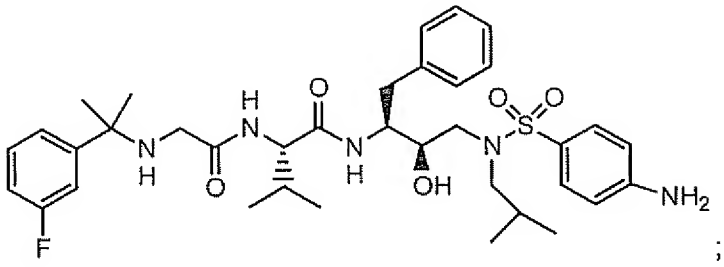

;

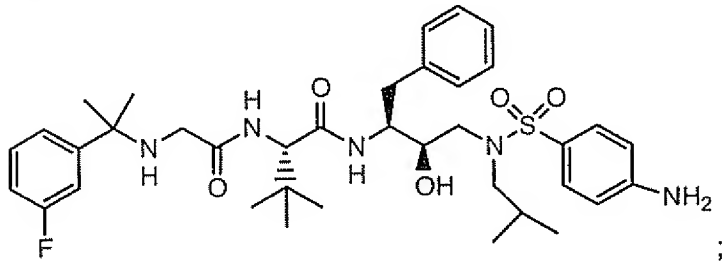

;

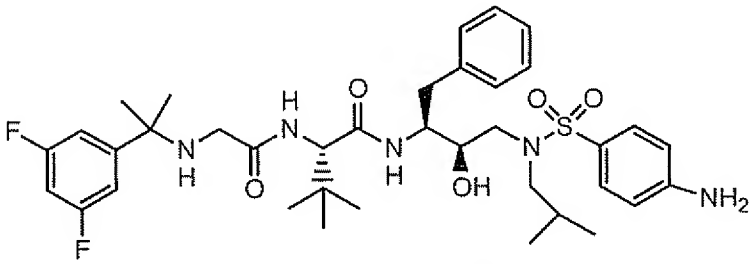

;

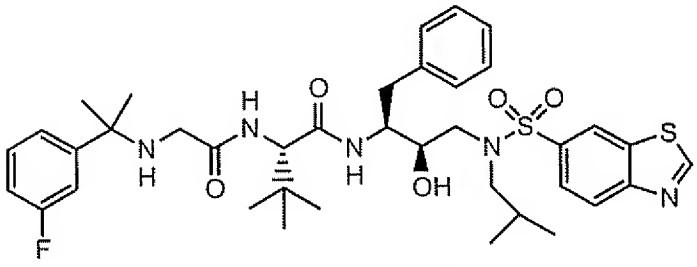

;

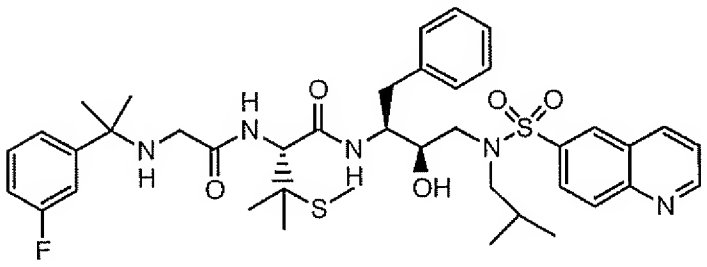

;

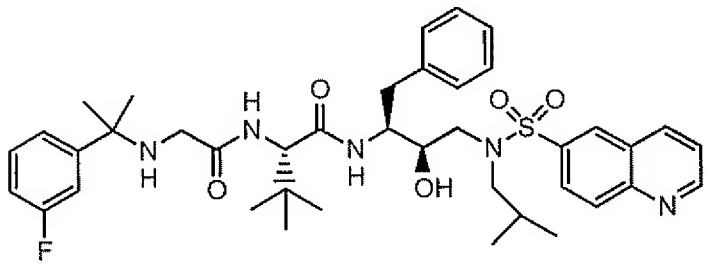

;

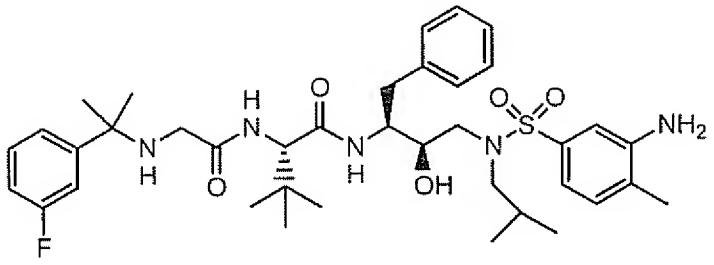

;

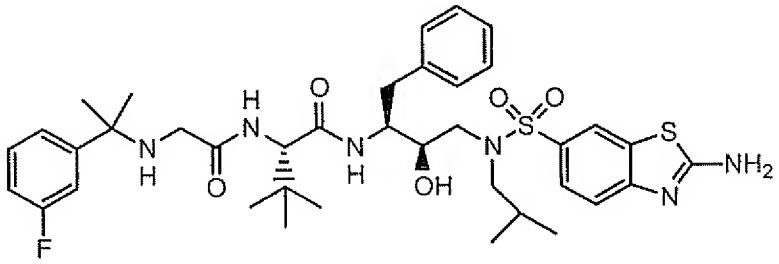

;

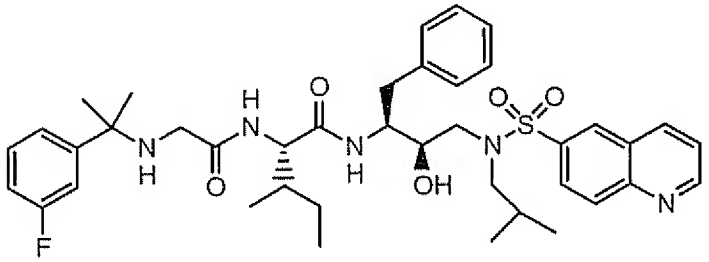

;

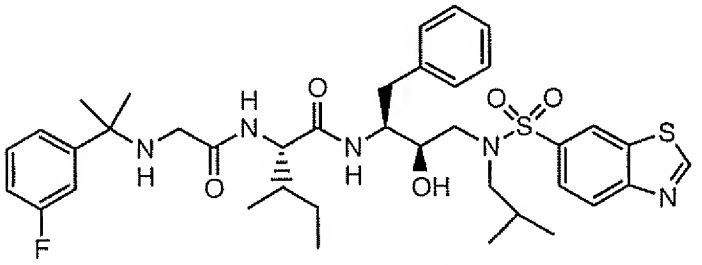

;

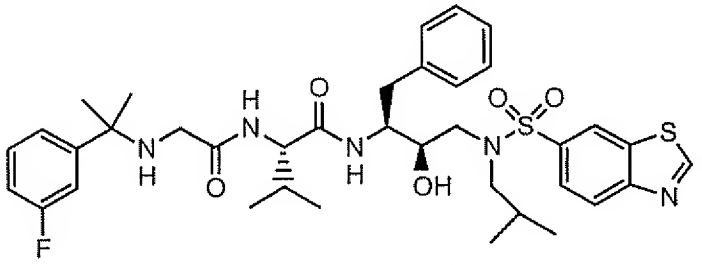

;

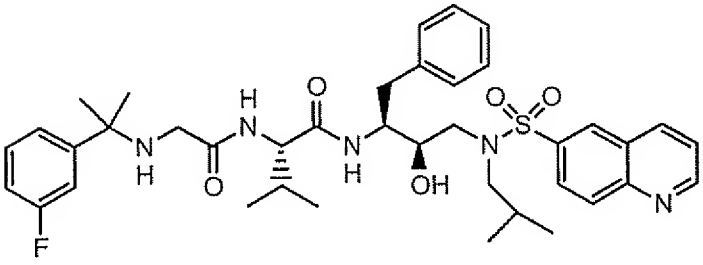

;

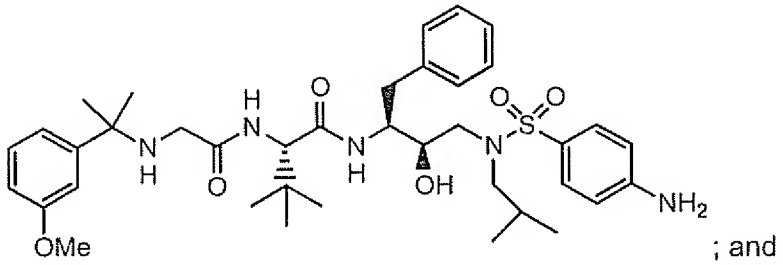

;


;

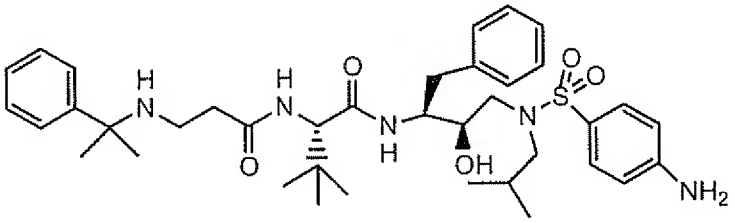

;


;


;



; and



or a pharmaceutically acceptable salt thereof.

11. (original) A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1 and a pharmaceutically acceptable carrier.

12. (canceled)

13. (original) A method for treating HIV infection comprising administering a therapeutically effective amount of a compound of Claim 1 to a patient in need of such treatment.

14. (original) The method of claim 13 comprising co-administering a therapeutic amount of an HIV reverse transcriptase inhibitor, an HIV protease inhibitor, or a combination thereof.

15. (original) A method for treating AIDS or ARC comprising administering a therapeutically effective amount of a compound of Claim 1 to a patient in need of such treatment.

16. (original) The method of claim 15 comprising co-administering a therapeutic amount of an HIV reverse transcriptase inhibitor, an HIV protease inhibitor, or a combination thereof.